Centers for Disease Control and Prevention

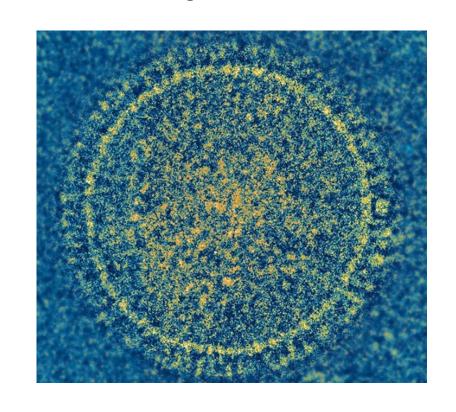




Introduction to ACIP's Maternal/Pediatric RSV Work Group

Sarah S. Long, MD
Chair, Maternal/Pediatric RSV Work Group

ACIP General Meeting January 12, 2022



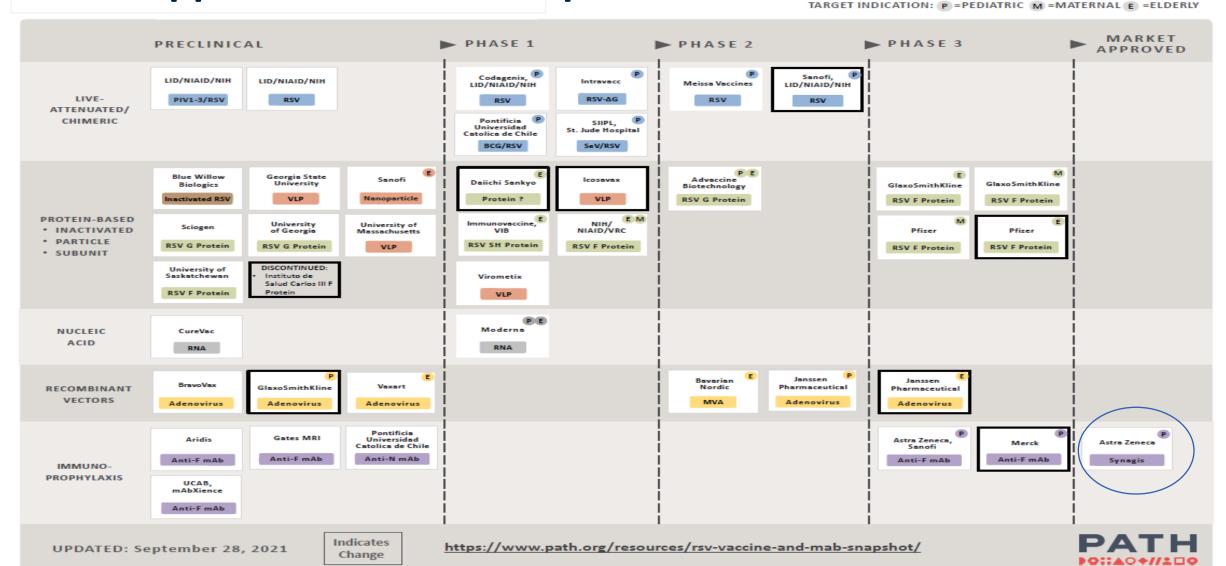
Purpose of the Work Group

- Respiratory syncytial virus (RSV) is a major cause of lower respiratory illness, particularly among infants and children and among older adults and adults with chronic medical conditions.
- RSV vaccine and monoclonal antibody development has progressed in the past decade with over 40 candidate vaccines and monoclonal antibodies currently in development.
- Target populations for whom these products are intended include infants and young children, pregnant women, and older adults.

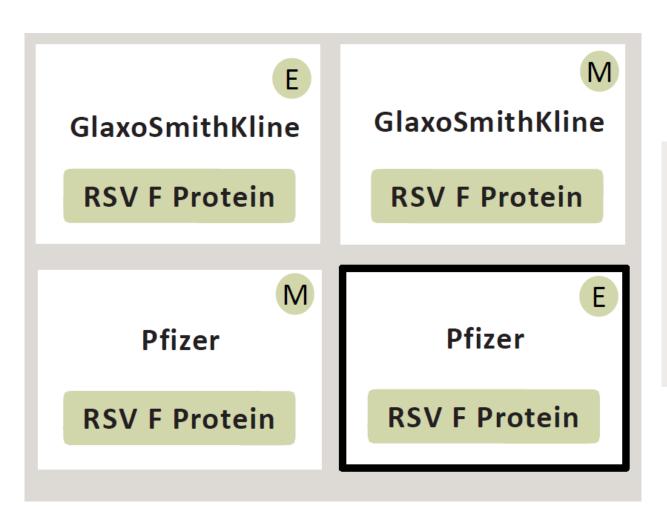
Work Group Activities

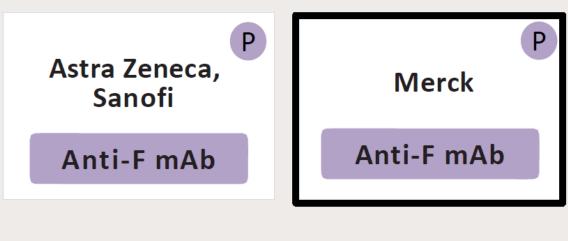
- Consider recommendations for use of RSV vaccines and monoclonal antibodies targeting protection of children aged <18 years:
 - Review the epidemiology and burden of RSV disease in children and pregnant women
 - Review efficacy, immunogenicity, safety and cost-effectiveness of RSV vaccine(s) and newly developed monoclonal antibody products in pregnant women and children
 - Provide evidence-based recommendations regarding use of RSV vaccine(s) and newly developed monoclonal antibody products in pregnant women and children
 - Identify areas in need of further research for informing potential future vaccine and monoclonal antibody recommendations.

RSV Vaccine and Monoclonal Antibody Products that are FDA-Approved or in Development



RSV Vaccine and Monoclonal Antibody Products in Phase 3 Trials





P = PEDIATRIC M = MATERNAL E = ELDERLY

Work Group Members

ACIP Members	Consultants
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Sarah Long (chair) Denise Jamieson (Emory University School of Medicine)

Pablo Sanchez Cody Meissner (Tufts Medical Center)

Oliver Brooks Helen Chu (University of Washington)

Ex Officio Members Daniel Feikin (World Health Organization)

Rachel Zhang (FDA-CBER)

Liaisons

CDC staff: Jefferson Jones (lead)

Jaya Goswami (FDA-CBER) Carol Baker (IDSA) Respiratory Viral Immunization Safety

Judy Beeler (FDA-CBER) Patsy Stinchfield (NFID) <u>Branch</u> <u>Office</u>

Yodit Belew (FDA-CDER) Brenna L. Hughes (ACOG) Angie Campbell Christine Olson

Prabha Viswanathan (FDA-CDER) Nicole Chaisson (AAFP) Meredith McMorrow Anne Hause

Samer El-Kamary (FDA-CDER) Sean O-Leary (AAP) Mila Prill Immunizations Service

Sonnie Kim (NIH-NIAID) Fiona Havers <u>Division</u>

April Killikelly (Public Health Natalie Thornburg Patricia Wodi

Agency of Canada)

Adam Macneil

Neil Murthy

Work Group Calls

- Held in October 2021–January 2022
 - Reviewed burden and epidemiology of RSV
 - Reviewed U.S. RSV surveillance systems and definitions of RSV infection and disease
 - Discussed potential components of cost effectiveness considerations for RSV vaccines and mAb products

Upcoming Work Group Activities

- Review safety and efficacy with manufacturers of vaccines and mAb
- Review RSV seasonality data in relation to potential recommendations for timing of administration of vaccines and mAb product
- Review results of cost effectiveness model
- Present data to ACIP
- Apply Evidence to Recommendation Framework for each product

The first vote by ACIP is not expected until 2023.

For more information, contact CDC 1-800-CDC-INFO (232-4636)
TTY: 1-888-232-6348 www.cdc.gov

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.

